

**DIPHENHYDRAMINE HCL- diphenhydramine hcl capsule**  
**Akron Pharma Inc.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Akron Pharma, Inc.**

***Active ingredient (in each banded capsule)***

Diphenhydramine Hydrochloride 25 mg

***Purpose***

Antihistamine

***Uses***

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

***Warnings***

***Do not use***

- to make a child sleepy
- with any other product containing diphenhydramine, even one used on skin

***Ask a doctor before use if you have***

- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis

***Ask a doctor or pharmacist before use if you are***

taking sedatives or tranquilizers.

***When using this product***

- marked drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- excitability may occur, especially in children
- be careful when driving a motor vehicle or operating machinery

**If pregnant or breast-feeding,**

ask a health professional before use.

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

**Directions**

- take every 4 to 6 hours
- do not take more than 6 doses in 24 hours

<b>Age</b>	<b>Dose</b>
adults and children 12 years and over	1 to 2 capsules
children 6 to under 12 years of age	1 capsule
children under 6 years of age	do not use this product in children under 6 years of age

**Other information**

- store at temperature 15° to 30° C (59° to 86°F)
- do not use if either capsule band or imprinted safety seal under cap is broken or missing
- protect from moisture

**Inactive ingredients**

colloidal silicon dioxide, croscarmellose sodium, hard gelatin capsules, hydroxypropyl methylcellulose, magnesium stearate, microcrystalline cellulose, propyl paraben sodium.

**Questions or comments?**

**Call toll-free 1-877-225-6999**

**Mfg Lic .No: TN/DRUGS/558/1997**

**Manufactured for:**

Akron Pharma, Inc,

373 RT US 46 W, Building E,

Suite 117, Fairfield, NJ 07004

Akron Pharma NDC 71399-8028-1

\*Compare to active ingredient in BENADRYL® Allergy

# Diphenhydramine HCl Capsules

## 25 mg

### Antihistamine

Complete Allergy Medication

For the temporary relief of the symptoms of  
 •Upper Respiratory Allergies •Hay Fever

**100 CAPSULES**

**Drug Facts**

**Active ingredient (in each banded capsule)**  
Diphenhydramine Hydrochloride 25 mg ..... Antihistamine

**Uses**  
 ■ temporarily relieves these symptoms due to hay fever or other upper respiratory allergies & common cold:  
 ■ runny nose ■ sneezing ■ itching of the nose or throat  
 ■ itchy, watery eyes

**Warnings**  
**Do not use** ■ to make a child sleepy ■ with any other product containing diphenhydramine, even one used on skin

**Ask a doctor before use if you have**  
 ■ glaucoma  
 ■ trouble urinating due to an enlarged prostate gland  
 ■ a breathing problem such as emphysema or chronic bronchitis

**Ask a doctor or pharmacist before use if you are**  
 taking sedatives or tranquilizers

**When using this product**  
 ■ marked drowsiness may occur  
 ■ avoid alcoholic drinks  
 ■ alcohol, sedatives and tranquilizers may increase drowsiness  
 ■ be careful when driving a motor vehicle or operating machinery  
 ■ excitability may occur, especially in children

**If pregnant or breast-feeding,**  
 ask a health professional before use.

**Keep out of reach of children.**  
 In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Child-Resistant Packaging**      **Drug Facts (Continued under label)**

\*This Product is not manufactured or distributed by Johnson & Johnson Consumer Inc., owner of the registered trademark Benadryl®.

Batch No. :  
Mfg. Date :  
Exp. Date :

**PEEL HERE FOR MORE DRUG FACTS**

Akron Pharma  
 Manufactured for:  
 Johnson & Johnson  
 373 RT US 46 W Building E  
 Suite 117, Fairfield, NJ 07004  
 www.akronpharma.com

**Drug Facts (continued)**

**Directions** ■ take every 4 to 6 hours ■ do not take more than 6 doses in 24 hours  
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**Mfg Lic. No: TN/DRUGS/558/1997**

274/MNC/US/101

<b>DIPHENHYDRAMINE HCL</b>			
diphenhydramine hcl capsule			
Product Information			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71399-8028
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg	
Inactive Ingredients			
<b>Ingredient Name</b>	<b>Strength</b>		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)			
GELATIN (UNII: 2G86QN327L)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			

<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>PROPYLPARABEN SODIUM</b> (UNII: 625NNB0G9N)	

### Product Characteristics

<b>Color</b>	pink (PINK WHITE)	<b>Score</b>	no score
<b>Shape</b>	capsule	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	AP;29
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-8028-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2021	
2	NDC:71399-8028-2	1000 in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2021	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	01/15/2021	

**Labeler** - Akron Pharma Inc. (067878881)

Revised: 2/2023

Akron Pharma Inc.